

In the Claims:

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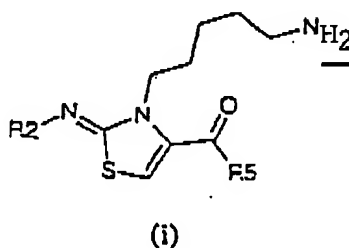
Claims 1 and 7 (cancelled).

Claim 8 (previously presented) A composition for an affinity for somatostatin receptors comprising an amount of a compound of Claim 12 sufficient for said affinity and a pharmaceutical carrier.

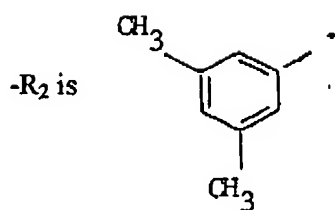
Cancel Claim 9.

Claims 10 and 11 (cancelled).

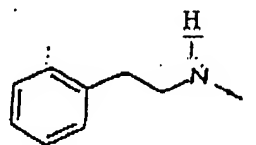
Claim 12. (currently amended) A compound of the formula:

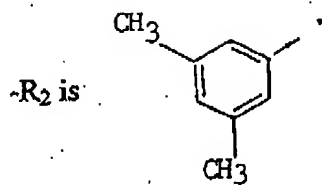
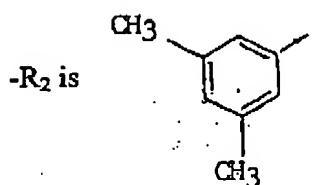
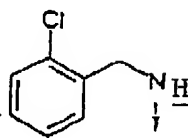
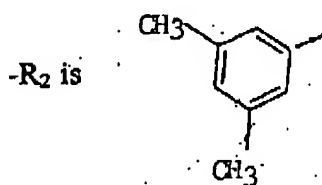
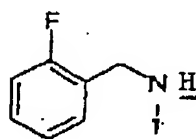
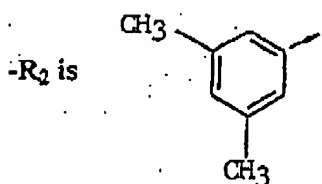
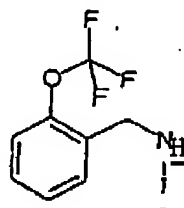
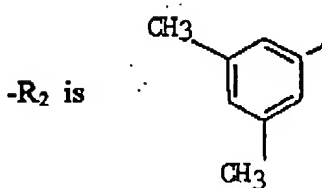
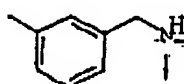
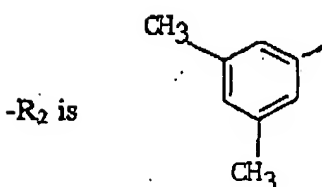
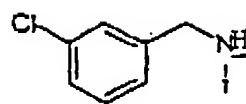
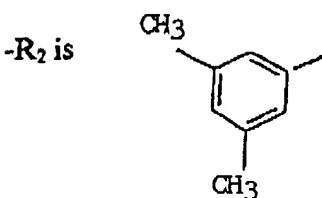
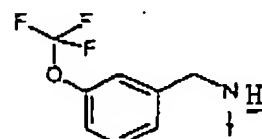
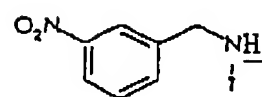


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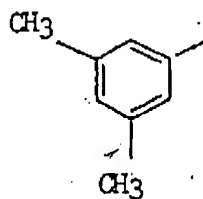


and  $R_5$  is

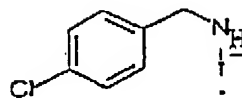


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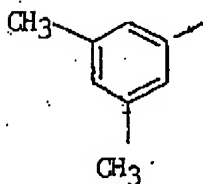
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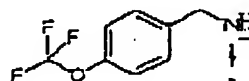
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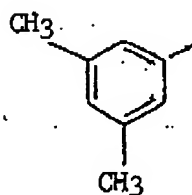
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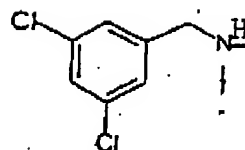
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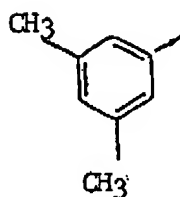
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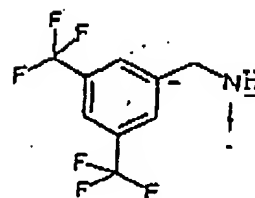
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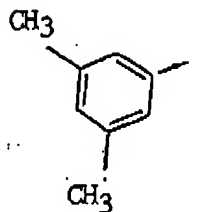
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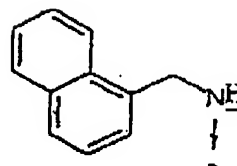
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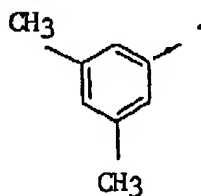
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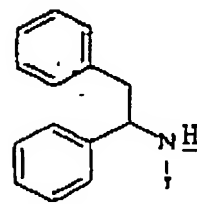
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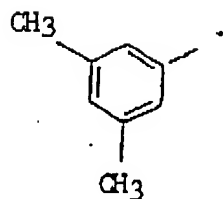
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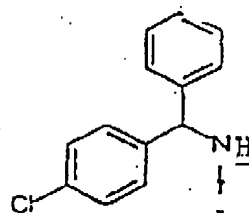
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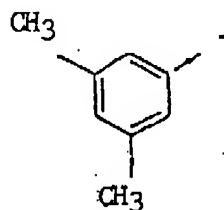
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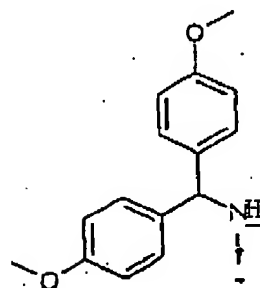
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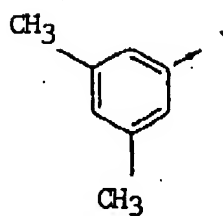
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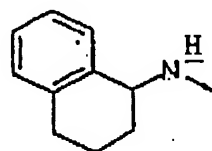
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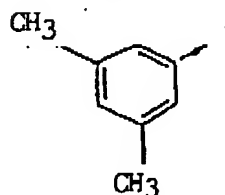
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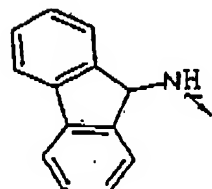
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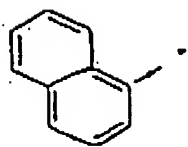
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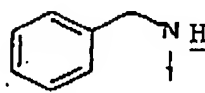
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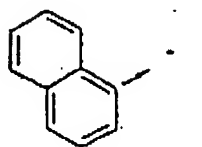
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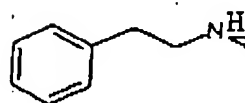
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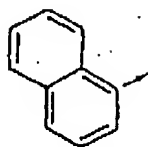
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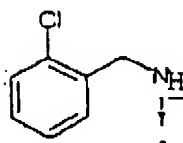
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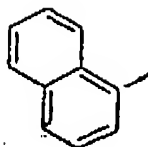
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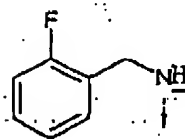
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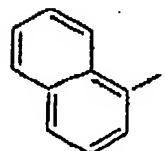
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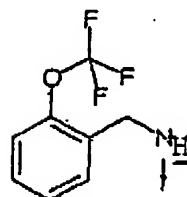
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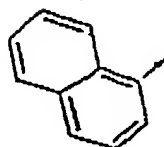
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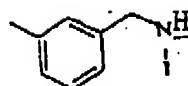
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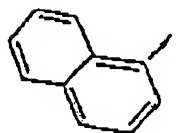
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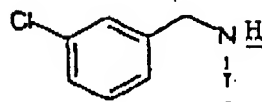
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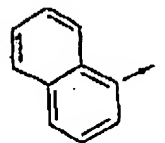
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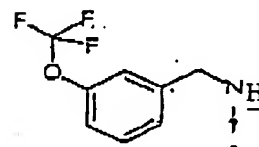
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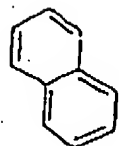
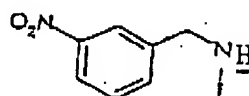
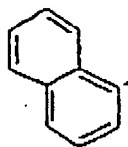
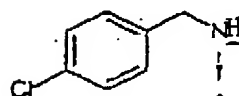
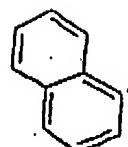
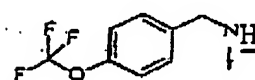
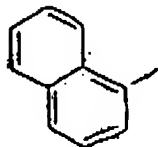
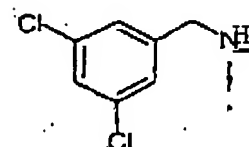
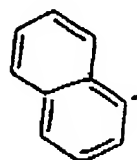
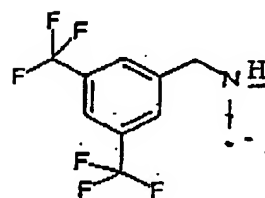
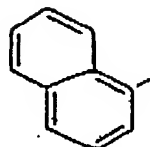
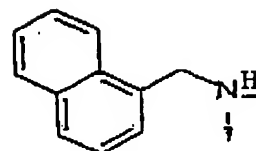
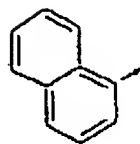
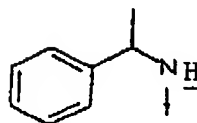


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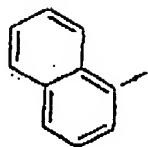


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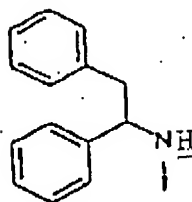


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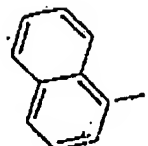
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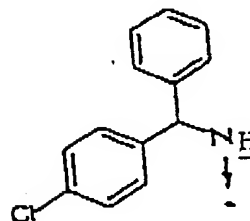
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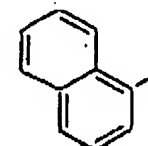
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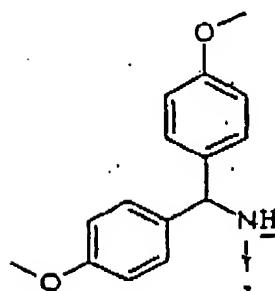
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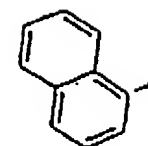
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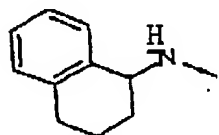
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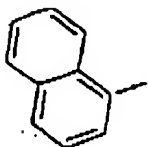


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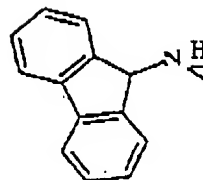


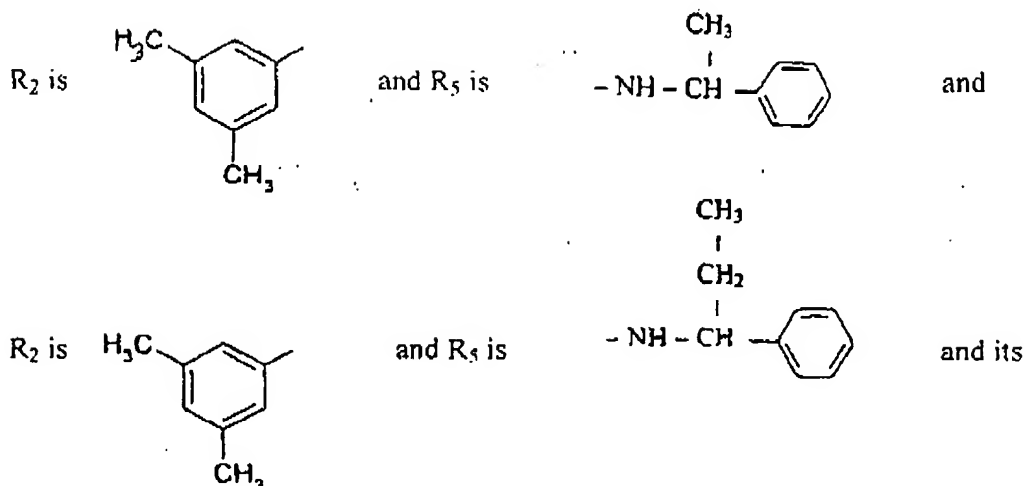
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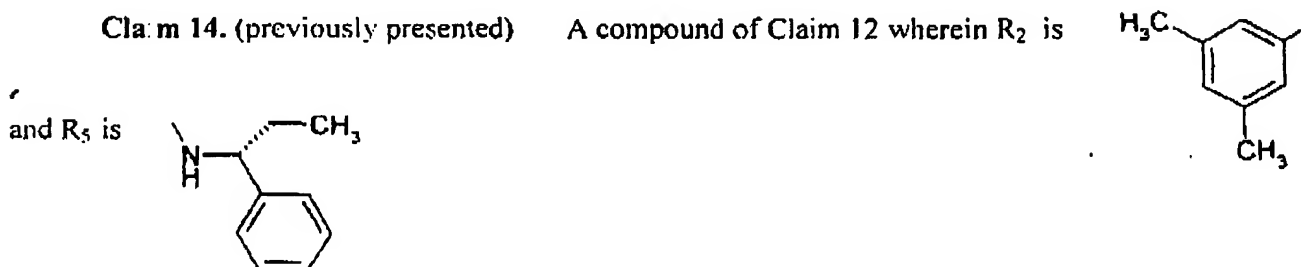
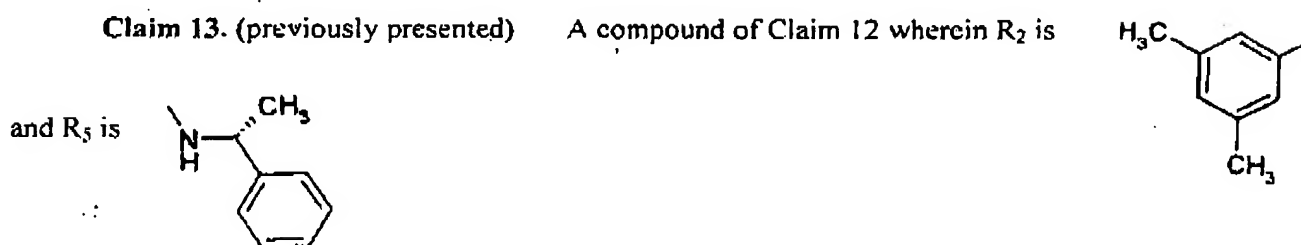


and R<sub>5</sub> is





it being understood that the nitrogen atom in formula (I) could be  $NH_2$  or a nitrile and that the nitrogen atom in the  $R_5$  groups is  $NH$  and its pharmaceutically acceptable salt.





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